

This Page Is Inserted by IFW Operations  
and is not a part of the Official Record

## **BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning documents *will not* correct images,  
please do not report the images to the  
Image Problem Mailbox.**

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
19 April 2001 (19.04.2001)

PCT

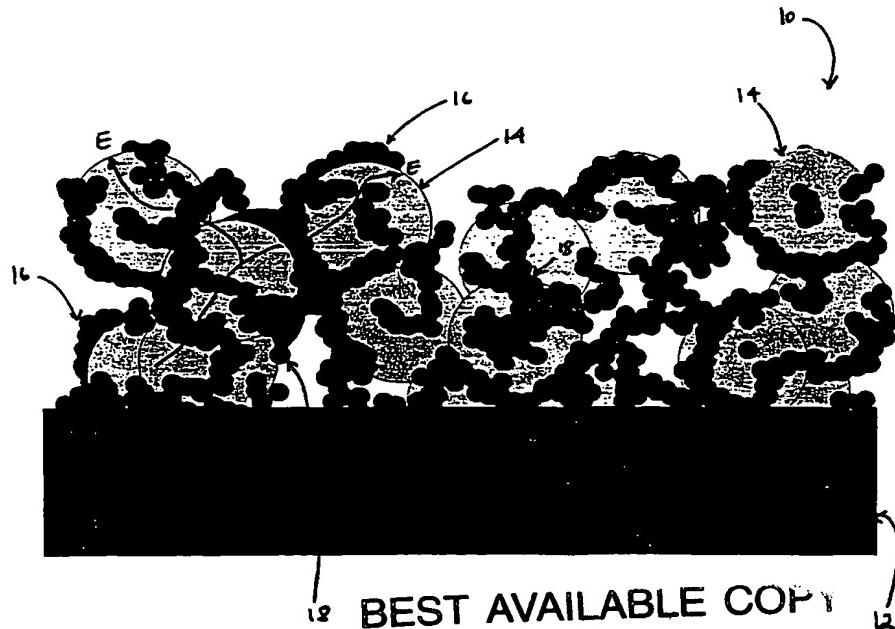
(10) International Publication Number  
**WO 01/28024 A1**

- (51) International Patent Classification<sup>7</sup>: H01M 8/12,  
4/86, G01N 27/56
- (21) International Application Number: PCT/CA00/01166
- (22) International Filing Date: 6 October 2000 (06.10.2000)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:  
60/158,124 8 October 1999 (08.10.1999) US  
60/231,542 11 September 2000 (11.09.2000) US
- (71) Applicant: GLOBAL THERMOELECTRIC INC.  
[CA/CA]; 4852 - 52 Street S.E., Calgary, Alberta T2B 3R2  
(CA).
- (72) Inventors: GHOSH, Debabrata; 4852 - 52 Street S.E.,  
Calgary, Alberta T2B 3R2 (CA). MARTELL, Frank;  
4852 - 52 Street S.E., Calgary, Alberta T2B 3R2 (CA).
- (74) Agent: BENNETT JONES LLP; 1000 Atco Centre,  
10035 - 105 Street, Edmonton, Alberta T5J 3T2 (CA).
- (81) Designated States (*national*): AE, AG, AL, AM, AT, AU,  
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ,  
DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,  
HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,  
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,  
NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,  
TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.
- (84) Designated States (*regional*): ARIPO patent (GH, GM,  
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian  
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European  
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,  
IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG,  
CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:  
— With international search report.

[Continued on next page]

(54) Title: COMPOSITE ELECTRODES FOR SOLID STATE ELECTROCHEMICAL DEVICES



WO 01/28024 A1

(57) Abstract: A solid state ionic device includes a dense electrolyte sandwiched between two porous electrodes. In one embodiment, the device is anode supported and the cathode is formed of a porous three-dimensional solid phase structure having an electrocatalytic phase of a plurality of electrocatalytic particles (14) and an ionic conducting phase of a plurality of ionic conductor particles (16). The mean or median size of the electrocatalytic particles is larger than the mean or median size of the ionic conductor particles. The device may further include a long range electronic conducting layer of lanthanum cobaltate or other electronically conducting material.



— *Before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments.*

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

## COMPOSITE ELECTRODES FOR SOLID STATE ELECTROCHEMICAL DEVICES

5

### CROSS-REFERENCE TO RELATED APPLICATIONS

This application claims the priority benefit of U.S. Provisional Application Nos. 60/158,124 filed on October 8, 1999 (Solid Oxide Fuel Cell Composite Electrode), and  
10 60/\_\_\_\_\_, filed September 9, 2000 (Improved Composite Electrodes For Solid State Devices), which applications are incorporated herein by reference in their entirety.

### FIELD OF THE INVENTION

15 The present invention relates to composite electrodes and functional layers for electrode reactions for use with solid-state ionic devices, and solid oxide fuel cells in particular.

### BACKGROUND OF THE INVENTION

20

The following references are referred to herein by their numerical reference and the contents of each is incorporated herein by reference.

1. Erning, J.W., Hauber, T., Stimming, U. Wippermann, K., Catalysis of the electrochemical processes on solid oxide fuel cell cathodes, Journal of Power Sources 61 (1996) 205-211.
2. M. Watanabe, H. Uchida, M. Shibata, N. Mochizuki and K. Amikura, High performance catalyzed -reaction layer for medium temperature operating solid oxide fuel cells, J. Electrochem. Soc., vol. 141, (1994) 342-346.

30

3. Sahibzada, M., Benson, S.J., Rudkin, R.A., Kilner, J.A., Pd-promoted  $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.2}\text{Fe}_{0.8}\text{O}_3$  cathodes. Solid State Ionics 113-115 (1998) 285-290.
4. M. M. Murphy, J. Van herle, A.J. McEvoy, K. Ravindranathan Thampi, Electroless deposition of electrodes in solid oxide fuel cells, J. Electrochem. Soc., vol. 141 (1994) 30 L94-96.
5. Uchida et al. Shin-ichi Arisaka and Masahiro Watanabe, Paper B-IN-05 at 121<sup>st</sup> International Conference on Solid State Ionics (1999) 154-155.

10

## Background

Solid state ionic devices typically consist of a fully dense electrolyte sandwiched between thin electrode layers. It is known that the principal losses in most solid state ionic devices occur in the electrodes or the electrode/electrolyte interfaces. Therefore, minimization of these losses is critical to efficient operation of these devices.

Solid oxide fuel cells (SOFC) are theoretically very efficient energy conversion devices that have the potential of becoming a commercial product for numerous uses. A SOFC is a solid electrochemical cell which consists of a solid electrolyte impervious to gases, sandwiched between a porous cathode and a porous anode. Oxygen gas is transported through the cathode to its interface with the electrolyte where it is reduced to oxygen ions, which migrate through the electrolyte to the anode. At the anode, the ionic oxygen reacts with fuels such as hydrogen or methane and release electrons. The electrons travel back to the cathode through an external circuit to generate electric power.

The construction of conventional SOFC electrodes are well known. Electrodes are often applied as composites of an electron conducting material and an ion conducting material. For instance, an anode may consist of electronic conducting nickel (Ni) and ionic conducting yttria

stabilized zirconia (YSZ) while the cathode may consist of a perovskite such as  $\text{La}_{1-x}\text{Sr}_x\text{MnO}_{3-\delta}$  (LSM) as the electron conducting material and YSZ as the ion conductor.

- Conventional SOFCs exhibit high performance at operating temperatures of 1000° C.
- 5 However, such high temperature operation has disadvantages such as physical or chemical degradation of the construction materials. Therefore, it is desirable to reduce the operating temperature of a SOFC stack to a medium temperature of about 700° C. However, at such medium temperatures, electrode reaction rates decrease significantly. Prior art efforts to increase electrode reactivity at lower temperatures have focussed on optimizing the electrode
- 10 microstructure and by introducing catalytic materials into the electrode structure.

It is well known to provide an activated surface on the fuel cell electrodes by means of a catalyst to aid the electrochemical process. Nickel is commonly used as a catalyst on the anode side for oxidation of fuel. On the cathode side, ceramic cathode materials typically used in

15 SOFCs, such as perovskites have a high activation energy for oxygen reduction. Therefore, the activation energy may be reduced for the oxygen reduction reaction by adding noble metals such as Au, Ag, Pt, Pd, Ir, Ru and other metals or alloys of the Pt group. Erning et al. [1] reported that addition of highly dispersed noble metals ( $\leq 0.1 \text{ mg/cm}^2$ ) lowers the activation energy of the oxygen reduction reaction at the cathode of an SOFC. M. Watanabe [2] also found that the

20 anodic polarization resistance and its activation energy were greatly decreased by loading only a small amount of catalyst such as Ru, Rh, and Pt onto a samaria-doped ceria (SDC) anode. A large depolarizing effect was also observed with a Pt-catalyzed LSM cathode, especially at high current densities. Sahibzada et al. [3] has recently reported that LSCF electrodes which were impregnated with small amounts of Pd resulted in 3-4 times lower cathodic impedance in the

25 temperature range 400 to 750°C. The overall cell resistance decreased 15% at 650°C and 40% at 550°C.

For economic reasons, noble metal catalysts are applied in very small amounts to catalyze the electrochemical process at electrodes. The catalysts are conventionally impregnated in the

pores of the electrode by a filtration or a chemical process. The impregnation process is frequently followed by a binding process where a binder is superimposed on the deposited particles to provide a secure and durable attachment of the coating with the base material. U.S. Patent Nos. 3,097,115; 3,097,974; 3,171,757 and 3,309,231 disclose such conventional  
5 impregnating processes for porous electrodes.

The catalysts may also be applied by common electroless deposition techniques for Ni, Pd and Ag [4] and replacement plating, as disclosed in U.S. Patent No. 3,787,244. In this process, an acidic plating solution containing a salt of a noble metal catalyst is forced through the  
10 pores of a nickel electrode substrate and the noble metal ions from the dissolved salt replace a thin layer of the nickel surface within the pores.

It is known [1] to form highly dispersed catalyst layers with an amount of less than 0. 1 mg/cm<sup>2</sup> from aqueous solutions of Pt, Pd, Ir or Ru salts. A few drops of these solutions were  
15 applied onto the electrolyte surface. After drying, the salts were either reduced to metal form by heating under hydrogen (Pt and Pd) or oxidized by heating under air (Ir and Ru). Most recently, Uchida et al. [5] applied nanometer-sized noble metal catalysts to both anode and cathode resulting in appreciably lower overpotential ohmic resistance.

20 Singheiser (EP 424813) discloses an intermetallic compound layer (0.5-5 µm) contains 2-70 wt.% of a noble metal such as Pt, Ag or Pd which can be used between electrolyte and electrodes, or to connect electrically two fuel-cells. It is claimed that the fuel cell can be operated at a lower temperature due to higher electrode conductivity.

25 Because of the cost of noble metals, the application of noble metals in SOFC electrodes so far are mainly limited to its catalytic abilities. All recent efforts have been to add very fine particles of the catalyst in order to maximize the three phase boundary of the catalyst, the gas phase and the electrolyte. The catalyst is either applied as a very thin layer at the electrolyte/electrode boundary or is widely dispersed throughout the electrode.

In U.S. Patent No. 5, 543,239 issued to Virkar et al., an electrocatalyst is incorporated into a electrode microstructure that is claimed to improve the performance of a solid state ionic device by providing a catalyst and by improving electrical conductance. In this disclosure, a  
5 porous ionic conductor is applied to a dense electrolyte substrate. An electrocatalyst is then introduced into the porous matrix to produce electrical continuity and a large three phase boundary line length. As a result, the electrocatalyst is applied as a thin layer of small particles over the ionic conductor.

10 The electrode disclosed by Virkar et al., however, does not solve the problem of electrode instability. It is known that vapor loss of noble metals occurs at even medium SOFC operating temperatures. According to the Thomson-Freundlich (Kelvin) equation, an important aspect of the vapor pressure difference across a curved surface is the increase in vapor pressure at a point of high surface curvature. Thus, the smaller the particle size, the higher the vapor pressure. This  
15 could cause significant vapor loss for small noble metal particles at SOFC operating temperatures.

Furthermore, higher vapor pressure at the particle surface and lower vapor pressure at a neck between two particles makes smaller particles much easier to be sintered. Thus, the  
20 microstructure of an electrode with submicronic noble metal (<0.5  $\mu\text{m}$ ) particles is not stable at medium to high SOFC operating temperatures, and especially when the electrode handles high current.

Furthermore, a thin electronic conducting layer at the electrode will have large ohmic  
25 resistance at the electrode which limits the current carrying capacity of the electrode. As shown in the current-voltage curves of the Virkar et al. patent, the experimental current is limited to 0.5 A/cm<sup>2</sup> for the Pt/YSZ and LSM/YSZ cathodes disclosed therein.

Therefore, there is a need in the art for a composite electrode which mitigates the

limitations of the prior art, allowing higher performance solid state ionic devices and solid oxide fuel cells in particular.

### **SUMMARY OF THE INVENTION**

5       The present invention is directed at an electrode having an improved microstructure which achieves a high density of active electrochemical reaction sites between the electrolyte and electrode and incorporates electrocatalytic materials such as noble metals into the electrode in an intimate fashion. As well, the improved microstructure may have better long-term structural  
10      stability of the cell by reducing the effects of noble metal catalyst sintering and vapor loss. The electrode may be incorporated into any solid state electrochemical devices such as oxygen pumps, membranes and sensors, solid state batteries or solid oxide fuel cells. The electrode of the present invention may be either a cathode or an anode.

15      Accordingly, in one aspect of the invention, the invention comprises an electrode forming part of a solid state electrochemical device, said electrode bonded to a dense electrolyte layer and comprising a porous three-dimensional solid phase comprising:

- (a)     an electrocatalytic phase comprising a plurality of electrocatalytic particles;
- 20       (b)    an ionic conducting phase comprising a plurality of ionic conductor particles;

25      wherein said electrocatalytic conducting phase and ionic conducting phase are interspersed and wherein the mean size of said electrocatalytic particles is substantially equal to or larger than the mean size of said ionic conducting particles.

The electrode of the present invention is formed by mixing ceramic ion conductor particles and noble metal electrocatalyst particles into a composite electrode which is then applied to a dense electrolyte substrate by screen printing or by similar well-known methods. The

- resulting electrode microstructure is highly porous and includes very long three-phase boundaries, direct ion conducting channels from the catalytic sites to the electrolyte and direct electron conducting channels through the electrode to the catalytic sites. The electrocatalyst particles are preferably comprised of a noble metal and are preferably larger than the ion conductor particles which results in a morphology where the ion conductor particles pin the boundaries of the noble metal particles. The relatively large noble metal particle size reduces vapor loss at elevated temperatures while grain boundary pinning reduces or prevents sintering or coalescing of the noble metal particles.
- 10 In one embodiment, the ion conductor particles may comprise ceramic particles which may preferably be yttrium stabilized zirconia and the noble metal particles may comprise palladium. Those skilled in the art will be aware of other materials which will function as ion conducting particles or as electrocatalytic particles.
- 15 In one embodiment, the invention may comprise an electrode comprising (a) an electrode functional layer for use in a solid state electrochemical device, said layer comprising a porous three dimensional structure comprising linked particles of an electrocatalytic material and linked particles of an ionic conductor wherein the median size of the electrocatalyst particles is approximately equal to or larger than the median size of the ion conducting particles; and (b) a long range conducting electrode layer that is applied on top of the functional layer. In a planar SOFC, long range conductivity refers to horizontal conductivity between the ribs of an interconnect plate, rather than the short range vertical conducting path through the functional layer. The conducting layer may comprise electronically conductive metal oxides such as lanthanum cobaltate.
- 20 In another aspect, the invention comprises a solid state electrochemical device comprising a porous anode, a dense electrolyte and a cathode comprising a porous three-dimensional structure comprising linked particles of an electrocatalytic material and linked particles of an ionic conductor wherein the mean or median size of the electrocatalyst particles is larger than the

mean or median size of the ion conducting particles. The solid state electrochemical device may be a solid oxide fuel cell.

In another aspect of the invention, the invention is a method of forming an electrode for  
5 use in a solid state electrochemical device having a dense electrolyte layer comprising the steps  
of:

- (a) mixing electrocatalytic particles with ion conducting particles where mean or median size of the electrocatalytic particles is substantially equal to or larger than the mean or median size of the ion conducting particles; and
- 10 (b) creating a porous three-dimensional structure bonded to the dense electrolyte layer, said structure comprising linked particles of the noble metal particles and linked particles of the ionic conductor.

In one embodiment, a further conducting layer of a metal oxide may be applied but not  
15 presintered to the electrode. The metal oxide may comprise lanthanum cobalt oxide.

#### **BRIEF DESCRIPTION OF THE DRAWINGS**

The invention will now be described by way of exemplary embodiments with reference to  
20 the accompanying drawings. In the drawings:

Figure 1 is a schematic representation of a cross-sectional view of a cathode according to one embodiment of the present invention.

25 Figure 2 is a scanning electron micrograph of a cathode cross-section according to one embodiment of the present invention.

Figure 3 is a graph of the I-V characteristics of a single fuel cell of the present invention.

Figure 4 is a scanning electron micrograph (5000x) of a cathode cross-section of another embodiment of the invention.

5       Figure 5 is a scanning electron micrograph of an alternative embodiment of a cathode in cross-section.

Figure 6 is a schematic depiction of current flow through the cathode and the conducting electrode.

10      Figure 7 is a graph of the I-V characteristics of one alternative embodiment of a single fuel cell.

Figure 8 is a graph of the I-V characteristic of one embodiment of a 15 fuel cell stack.

15      Figure 9 shows the effects of altering the noble metal concentration on power density, with tests conducted at different temperatures.

#### DETAILED DESCRIPTION OF THE INVENTION

20      The present invention provides for a composite electrode for use with a solid oxide fuel cell and further provides for a method of making such an electrode. When describing the present invention, all terms not specifically defined herein have their common art-recognized meanings.

##### A.     Definitions

25

As used herein, the term "about" refers to a range of values that is the stated value plus or minus 10%.

As used herein, the term "electrocatalyst" refers to a material which is both electronically

conducting and a catalyst for an electrode reaction. Electrocatalyst materials may include noble metals and certain metal oxides.

As used herein, the term "noble metal" refers to metals and alloys of the group  
5 comprising silver, gold, iridium, osmium, palladium, ruthenium, rhodium and platinum.

As used herein, the term "LC" or "lanthanum cobaltate" refers to LaCoO<sub>3</sub>.

#### B. Description

10 As shown in Figure 1, one embodiment of a porous composite electrode (10) is shown bonded to an electrolyte (12). The composite electrode is formed from electrocatalytic noble metal particles (14), and from ion conducting ceramic particles (16) which are bonded intimately to the electrolyte (12). The ceramic particles combine to form ionic conducting paths (I) from the  
15 electrolyte (12) to the electrochemical active sites (18). The metal phase forms electronic conducting paths (E) through the electrode (10) to the contact paste (not shown) and cathode electronical conducting strip (not shown). The electrochemical active area coincides with the three phase boundary (18) which extends along the common boundary of the gaseous pore phase, the ceramic phase (16) and the noble metal phase (14). It is generally believed that the electrode  
20 reaction substantially takes place at this boundary, where the three phases (gas, electrocatalyst conductor and ion conductor) meet.

Thus, the composite electrode of the present invention may provide more electrode  
reaction sites to lower the overpotential loss. Furthermore, the presence of catalytic noble metals  
25 at the electrochemical active areas (18) lowers the activation energy for the electrode reactions.

The ceramic ionic conducting phase in the composite electrode may be any known ion conductor such as yttria stabilized zirconia (YSZ). In a preferred embodiment, the ceramic phase is preferably the same material as the electrolyte so that interface between the ceramic phase and

the electrolyte is chemically stable and there is a good thermal match between the two materials.

The electrocatalytic phase may be any noble metal or noble metal alloy. These metals all have catalytic effect for the reduction of oxygen and are good electronic conductors. In a

5 preferred embodiment, palladium is used because its coefficient of thermal expansion is similar to that of the YSZ which may be used as the electrolyte and in the ceramic phase. Accordingly, the use of palladium and YSZ in the preferred composite electrode of the present invention provide good thermal stability even where the electrode is subjected to thermal cycling.

10 The relative proportion of the noble metal and ceramic ionic conducting phases may be varied. However, if the volume percentage of one phase is lowered too far, continuous channels of that phase may not form when the electrode is formed. It is preferable to have continuous ionic conducting channels, electronic conducting channels and porous channels throughout the composite electrode thickness.

15

The electronic conducting channels lowers the ohmic resistance of the cell. Electronic conductivity of the composite electrode may be increased by increasing the particle size of the noble metals and by increasing the volume percentage of the metal phase. However, increasing the particle size decreases the catalytic effect of the electrocatalyst. Ionic conductivity may be  
20 increased by decreasing the particle size of the ceramic material and by increasing the volume percentage of the ceramic phase. However, a longer three phase boundary is created by using smaller particles of either the ceramic or metal phase.

As shown in the Figures, because the ceramic particles are preferably smaller than the  
25 metal particles, the ceramic particles (16) partially cover the noble metal particles (14). This reduction in surface area of the metal phase reduces vapor loss of the noble metal at elevated operating temperature. Moreover, the ceramic particles (16) tend to agglomerate between two adjoining metal particles (14), in an effect known as grain boundary pinning, which prevents further sintering of noble metal particles. Thus, the morphology of the electrode, the

electrode/electrolyte interface and the three phase boundary may be more stable.

In one embodiment illustrated in Figure 1, the gas phase, the metal phase and the ceramic phase are approximately equal in volume percent. However, the metal particles are

5 approximately 5 to 10 times the size of the ceramic particles. The resulting microstructure is as shown in Figure 1 and Figure 2. As is apparent, the ceramic particles form continuous ion conducting channels in the form of particle chains to the electrolyte from the three phase boundary. The metal particles connect to form continuous electron conducting channels between the three phase boundary and the cathode conducting layer. Finally, the high porosity of the  
10 structure combined with the intertwining of the ion conducting channels and the electron conducting channels creates a tremendously large three phase boundary.

A feature of the present invention is the relative size of the metal particles compared to the ceramic particles. The metal particles should preferably be larger than the ceramic particles

15 and more preferably about 2 to 10 times larger. As a result of this size differential, the ceramic particles tend to agglomerate on the metal particles in continuous strings. In particular, the ceramic particles agglomerate along the contact patches of adjoining metal particles. As referred to above, this morphology not only increases the three phase boundary of the cathode but also reduces sintering of the metal particles and reduces evaporative loss of the metal.

20

An electrode according to the present invention may be applied to an electrolyte/anode substrate according to well known suitable techniques such as screen printing, tape casting, slip casting, vapor deposition or thermal spraying. A preferred method is screen printing using a paste formed from a suitable binder, a suitable solvent, the noble metal particles and the ion conductor  
25 particles. The nature and use of the binder and the solvent are well known to those skilled in the art.

In an alternative embodiment of the invention, a porous composite functional cathode layer (110) is shown bonded to an electrolyte (112) in Figures 4 and 5. The composite functional layer

(110) is formed from electron conducting and catalytic noble metal particles (114), and from ion conducting ceramic particles (116) which are bonded intimately to the electrolyte (112). Covering the functional layer (110) containing the catalytic noble metal particles (114) and the ion conducting ceramic particles (116) is a high electronically conductive layer (120). In one embodiment, the 5 electronically conductive layer (120) is made from LC material. Other suitable materials for use in an oxidizing environment may include LSM ( $\text{LaSrMnO}_3$ ), or other electronically conducting metal oxides.

In one embodiment the functional layer (110) comprises noble metal particles about 1  $\mu\text{m}$  10 in diameter and is about 1-5  $\mu\text{m}$  thick, and thus is about 1 to 5 particles thick. This provides a layer with good short range vertical electronical conductivity, since the probability of noble metal particles providing an electronical conducting path between the electrolyte and the LC layer (120) is greatly increased over the prior art, which features thicker layers and the same quantities of noble metals.

The ceramic particles are preferably smaller than the metal particles and may be in the range of 15 about 0.1 to 0.2  $\mu\text{m}$  in diameter.

In one embodiment, the electrode layer (10) or the noble metal functional layer (110) is comprised of 50% electrocatalytic particles and 50% ion conducting particles with about 33% porosity by volume. In other words, the electrode comprises 1/3 ion conducting particles, 1/3 20 electrocatalytic particles, and 1/3 pore space by volume. All references herein to volume percentage of the electrocatalytic phase is of the volume of the solid phase. This volume percentage of the electrocatalyst may be varied between about 1.0% and about 95% by volume of the solid portion of the electrode, and preferably between about 20% to about 60%, depending upon the cost target to be achieved, desired performance per cell, or other factors. The volume percentage of the electrode 25 taken by pore space is preferably about 30% or 1/3, although the electrode porosity may be higher or lower.

Where the electrocatalyst is a noble metal, the volume percentage of noble metal may be between about 1% to about 50% to achieve a good balance between cost and performance. As

shown in Figure 9, there is no increase in performance in cells having a noble metal content in the functional layer (110) higher than 50%, therefore the additional cost of the extra noble metal is not preferred. Where the highest performance is desired, the noble metal content is preferably about 50%. Where a balance of performance and cost is desired, the noble metal content is preferably about 5%. Although the cell performance decreases as the noble metal content is decreased towards 1%, the loss in performance may be offset by the reduced cost of manufacturing the cell.

In an embodiment where the noble metal layer is thin, less than about 5  $\mu\text{m}$ , and when the concentration of noble metals is below about 30 vol%, vertical conductivity (as indicated by arrow V in Figure 6) is much more probable than horizontal long range conductivity (H) because of the distribution of the noble metal particles. There may not enough metal particles to provide continuous long range (H) conducting paths to carry current to the relatively widely spaced ribs (130) of the interconnect plate (132). To overcome this difficulty, an electrode (120) of electronically conducting material is applied over the functional layer (110). This conducting electrode (120) may preferably be in the order of about 15 to about 20  $\mu\text{m}$  in thickness, but can vary from about 3 to about 100  $\mu\text{m}$ . The conducting electrode or "LC layer" (120) material is preferably lanthanum cobaltate ( $\text{LaCoO}_3$ ) which, in an oxidizing environment, has very good electronic conductivity properties although other suitable conductive materials may be used.

The LC layer is preferably not prefired prior to operation in the stack because it is preferred to avoid sintering of the LC layer. Once sintered, the LC layer has a thermal expansion rate about twice that of the remaining components in the fuel cell, with the resultant sealing and bonding problems due to thermal expansion mismatch. Also, LC can chemically react with YSZ forming undesirable phases at the high temperature encountered during sintering. For this reason, the LC layer is preferably not sintered prior to use within the fuel cell stack.

The combination of the thinner noble metal functional layer (110) and the long range electronic conducting LC layer (120) have produced a fuel cell that delivers improved performance over the prior art, and may deliver power densities in the region of 1.2 W/cm<sup>2</sup> as shown in Figure

9.

The following examples are intended to be illustrative of the claimed invention but not limiting thereof.

5    **Example 1**

This example discloses a method of making a Pd and YSZ composite cathode for an anode supported solid oxide fuel cell. The resulting cathode is schematically illustrated in Figures 1. A scanning electron micrograph of a cathode made in accordance with this example is  
10    shown in Figure 2.

A screen printable composite cathode paste was made up of equal volumes of well-dispersed Pd particles, 8 mole percent yttria stabilized zirconia (8YSZ) in alpha-terpineol. Ethyl-cellulose binder was added in an effective amount. The Pd particle size ranged from 0.5 to  
15    2  $\mu\text{m}$  with a median size of about 1  $\mu\text{m}$  while the 8YSZ particle size ranged from 0.1 to 0.2  $\mu\text{m}$  with a median size of about 0.17  $\mu\text{m}$ . The substrate (100 mm in square) consisted of a fully dense 8YSZ electrolyte (10  $\mu\text{m}$  thick) on a porous NiO-8YSZ anode (1 mm thick). The cathode paste was screen printed on the electrolyte side of the substrate. The foot prints were 90 mm in square. The prints were oven dried at 60 - 80°C, then fired at 1300°C in air for 2 hours. The  
20    thickness of the composite cathode after firing was about 5-10  $\mu\text{m}$ . The resulting solid phase was 50% vol Pd and 50% vol YSZ with approximately 33% porosity.

A comparison of the resulting Pd/8YSZ cathode cell with a similar cell with a common perovskite cathode (LSM) showed that the cell with Pd/8YSZ cathode had much better  
25    performance. A 15-cell stack made from cells with this composite cathode was tested at 750°C and achieved a power of 750 W with hydrogen/argon (50/50) mixture as the fuel. Current interrupt experiments showed that the improvement resulted from both lower ohmic resistance at the cathode due to the palladium conductivity network and lower overpotential loss due to an increase in the electrochemical active area (three-phase boundary) and catalytic active area

(palladium surface).

Figure 3 illustrates the I-V characteristics of a single fuel cell incorporating this embodiment of an electrode operating at temperatures varying from 600°C to 900°C.

5

### Example 2

10 This example discloses a Pd, YSZ, and LC composite cathode for an anode supported solid oxide fuel cell and a method of making such a cathode. A scanning electron micrograph of the resulting cathode is illustrated in Figure 4.

15 A screen printable composite cathode functional layer paste was made up of suitable volumes of well dispersed Pd particles and 8YSZ in alpha terpineol to achieve a solid phase of 5% Pd/95% 8YSZ. Ethyl cellulose binder was added in an effective amount. The Pd particle size ranged from 0.5 to 2 µm with a median size of about 1 µm while the 8YSZ particle size ranged from 0.1 µm to 0.2 µm with a median size of about 0.17 µm. The substrate (100mm square) consisted of a fully dense 8YSZ electrolyte (10 µm thick) on a porous NiO – 8YSZ anode (1 mm thick). The cathode functional layer paste was screen printed on the electrolyte side of the substrate. The footprints were 90 mm square. The prints were oven dried at 60-80 °C, then fired at 1300° C in air for 1 hour. The thickness of the composite functional layer after firing was about 1-3 µm. The LC layer was screen printed to a thickness of about 3 µm on top of the functional layer but was not sintered. Once the cell 20 is at operating temperature of 800 °C the LC powder bonded adequately to the functional layer.

25 Figure 7 illustrates the I-V characteristic of a single fuel cell incorporating this embodiment of a cathode at operating temperatures varying from 600°C to 900°C.

Figure 8 illustrates the I-V performance of a stack of 15 fuel cells incorporating this embodiment of a cathode.

Example 3

A composite cathode was screen printed in a similar manner as Example 2 above but to a thickness of about 10  $\mu\text{m}$ . The LC layer was again screen printed on top of the functional layer but to a thickness exceeding 30  $\mu\text{m}$ . A scanning electron micrograph showing the resulting cathode in cross-section is shown in Figure 5.

Example 4

10

Figure 9 illustrates the effect on power density ( $\text{w/cm}^2$  at 0.7v) of varying the proportion of palladium from 0% vol to 95% vol of the solid phase. As may be seen, performance is maximized with 50% vol Pd. However, significant performance is still achieved with Pd loading as low as 5% vol.

15

As will be apparent to those skilled in the art, various modifications, adaptations and variations of the foregoing specific disclosure can be made without departing from the scope of the invention claimed herein.

**WHAT IS CLAIMED IS:**

1. An electrode forming part of a solid state electrochemical device, said electrode bonded to a dense electrolyte layer and comprising a porous three-dimensional solid phase comprising:

5 (a) an electrocatalytic phase comprising a plurality of electrocatalytic particles;

10 (b) an ionic conducting phase comprising a plurality of ionic conductor particles;

15 wherein said electrocatalytic phase and ionic conducting phase are interspersed and wherein the mean size of said electrocatalytic particles is substantially equal to or larger than the mean size of said ionic conducting particles.

20 2. The electrode of claim 1 wherein the solid phase of the electrode is comprised of about 1% to about 95% electrocatalytic phase by volume.

25 3. The electrode of claim 2 wherein the solid phase of the electrode is comprised of about 1% to about 50% electrocatalytic phase by volume.

4. The electrode of claim 3 wherein the solid phase of the electrode is comprised of about 5% electrocatalytic phase by volume.

5. The electrode of claim 3 wherein the solid phase of the electrode is comprised of about 25 30% electrocatalytic phase by volume.

6. The electrode of claim 3 wherein the solid phase of the electrode is comprised of about 50% electrocatalytic phase by volume.

7. The electrode of claim 1 wherein the mean size of said electrocatalytic particles is at least about twice as large as the mean size of the ionic conductor particles.
8. The electrode of claim 7 wherein the mean size of said electrocatalytic particles is at least about four times as large as the mean size of the ionic conductor particles.  
5
9. The electrode of claim 1 wherein the electrode is a cathode and forms part of a solid oxide fuel cell comprising an electrolyte layer and an anode layer.
10. The electrode of claim 9 wherein the cathode is less than about 10 µm thick.  
10
11. The electrode of claim 10 wherein the cathode is less than about 5 µm thick.
12. The electrode of claim 11 wherein the cathode is less than about 3µm thick.  
15
13. The electrode of claim 1 wherein said electrode has a vertical dimension and a horizontal dimension and said electrocatalytic phase provides continuous electronical conductivity in the vertical dimension but not in the horizontal dimension.
20. 14. The electrode of claim 1 wherein said electrode further comprises a horizontally electronic conducting layer comprising an electronically conductive metal oxide.
15. The electrode of claim 14 wherein said electronically conductive metal oxide comprises lanthanum cobaltate.  
25
16. The electrode of claim 1 wherein the electrocatalytic particles are comprised of a noble metal.
17. The electrode of claim 16 wherein the noble metal is palladium.

18. The electrode of claim 1 wherein the ion conductor particles are comprised of YSZ.
19. A solid state electrochemical device comprising a porous anode, a dense electrolyte and a cathode comprising a porous three-dimensional structure comprising linked particles of an electrocatalytic material and linked particles of an ionic conductor wherein the mean or median size of the electrocatalyst particles is larger than the mean or median size of the ion conducting particles.  
5
- 10 20. The solid state electrochemical device of claim 19 wherein the device is a solid oxide fuel cell.
- 15 21. The fuel cell of claim 20 wherein the electrocatalytic material is a noble metal.
- 15 22. The fuel cell of claim 21 wherein the noble metal is palladium.
- 20 23. The fuel cell of claim 20 wherein the mean or median electrocatalytic particle size is at least about 2 times larger than the mean or median ion conductor particle size.
- 20 24. The fuel cell of claim 23 wherein the mean or median electrocatalytic particle size is about 4 to about 10 times larger than the mean or median ion conductor particle size.  
25
- 25 25. The fuel cell of claim 20 wherein the ionic conductor is comprised of the same ion conducting material as the electrolyte layer.
26. The fuel cell of claim 25 wherein the electrolyte and ion conducting particles are both comprised of YSZ.  
27. The fuel cell of claim 20 wherein the cathode is less than about 10  $\mu\text{m}$  thick.

28. The fuel cell of claim 27 wherein the cathode is less than about 5  $\mu\text{m}$  thick.
29. The fuel cell of claim 28 wherein the cathode is less than about 3  $\mu\text{m}$  thick.  
5
30. The fuel cell of claim 20 further comprising a long range electronically conductive layer applied to the cathode.
31. The fuel cell of claim 30 wherein said long range electronically conductive layer is comprised of lanthanum cobaltate.  
10
32. The fuel cell of claim 31 wherein said long range electronically conductive layer is not sintered prior to operation of the fuel cell.
- 15 33. A method of forming an electrode for use in a solid state electrochemical device having a dense electrolyte layer comprising the steps of:
  - (c) mixing electrocatalytic particles with ion conducting particles where mean or median size of the electrocatalytic particles is substantially equal to or larger than the mean or median size of the ion conducting particles; and  
20
  - (d) creating a porous three-dimensional structure bonded to the dense electrolyte layer, said structure comprising linked particles of the noble metal particles and linked particles of the ionic conductor.
- 25 34. The method of claim 33 wherein the electrocatalytic particles, the ion conducting particles , a suitable organic binder and a suitable solvent are mixed in appropriate volumes to form a paste which is then screen printed onto the dense electrolyte.
35. The method of claim 33 wherein the electrocatalytic particles are comprised of a noble

metal.

36. The method of claim 35 wherein the noble metal comprises palladium.
- 5 37. The method of claim 33 wherein the ion conducting particles are comprised of YSZ.
38. The method of claim 33 further comprising the steps of sintering the electrode and then applying a long range electronic conducting layer comprising a metal oxide.
- 10 39. The method of claim 38 wherein the metal oxide comprises lanthanum cobaltate.
40. The method of claim 33 wherein the volume percentage of electrocatalytic particles in the solid phase of the electrode is between about 1% and about 50%.

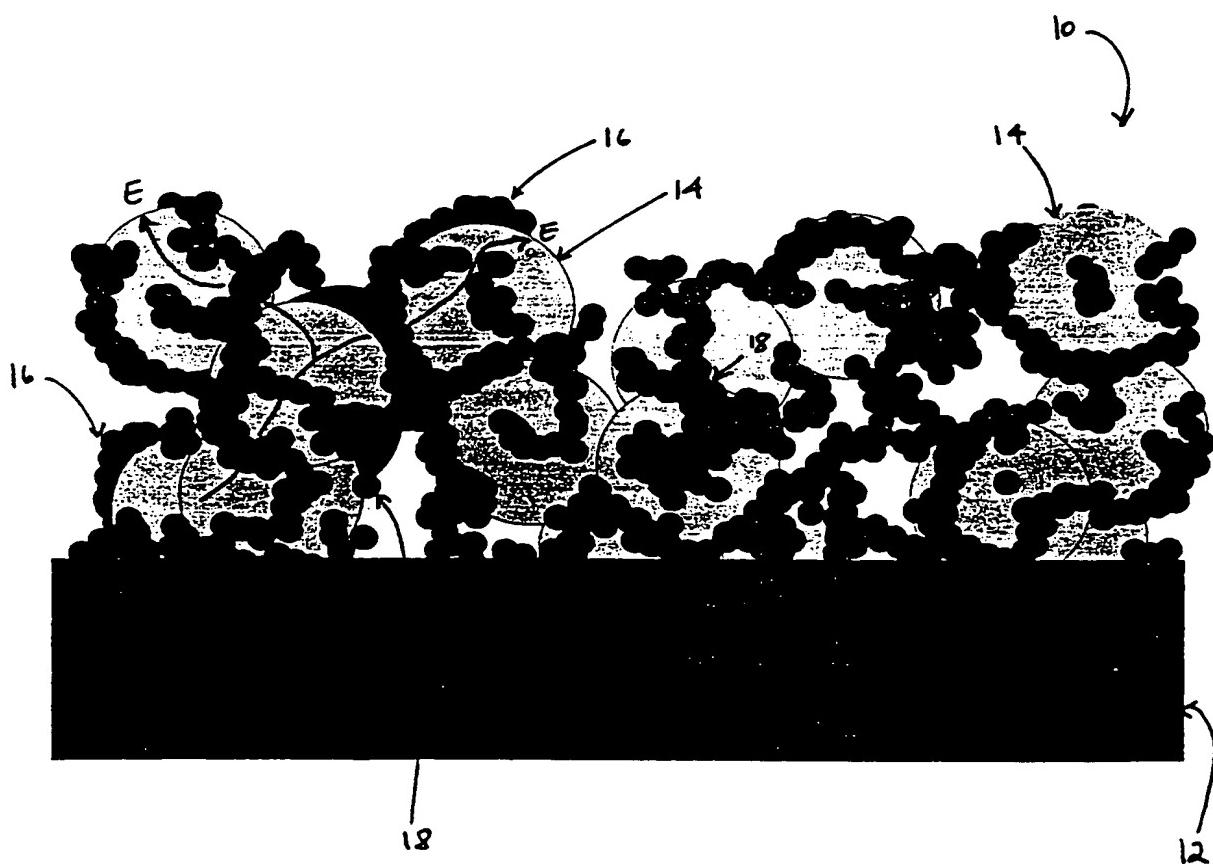


FIGURE 1

FIGURE 2

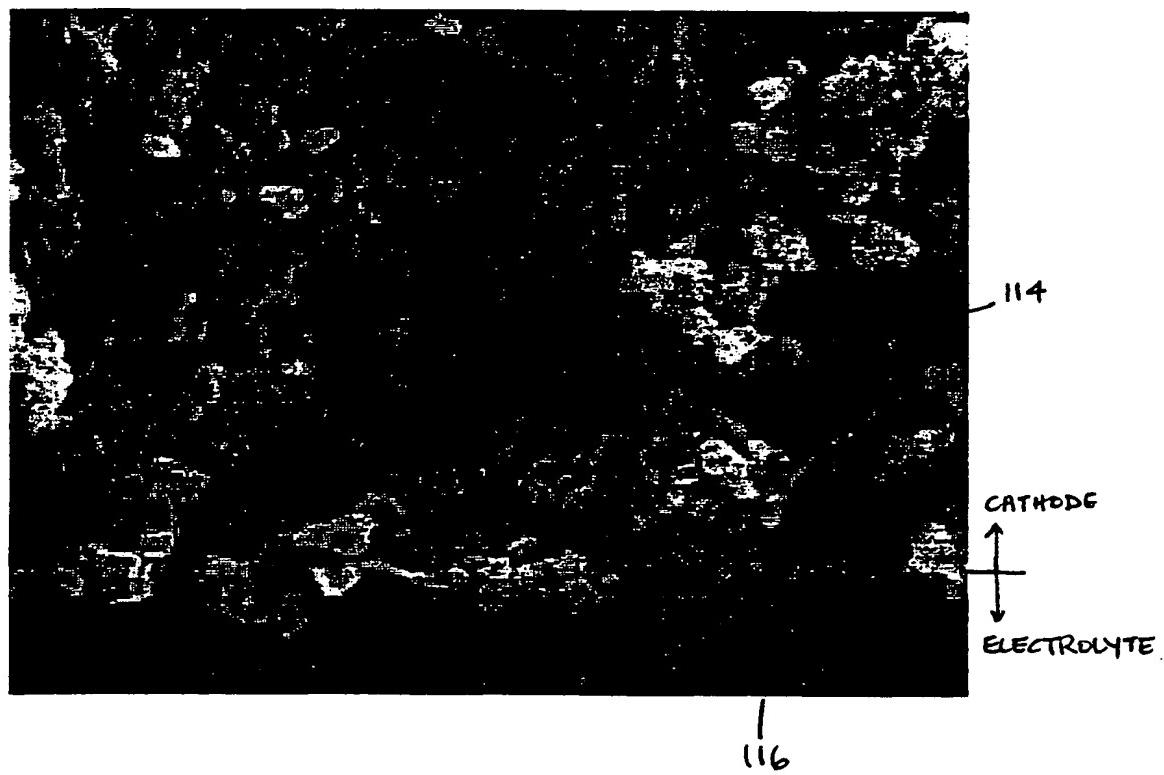


FIGURE 3

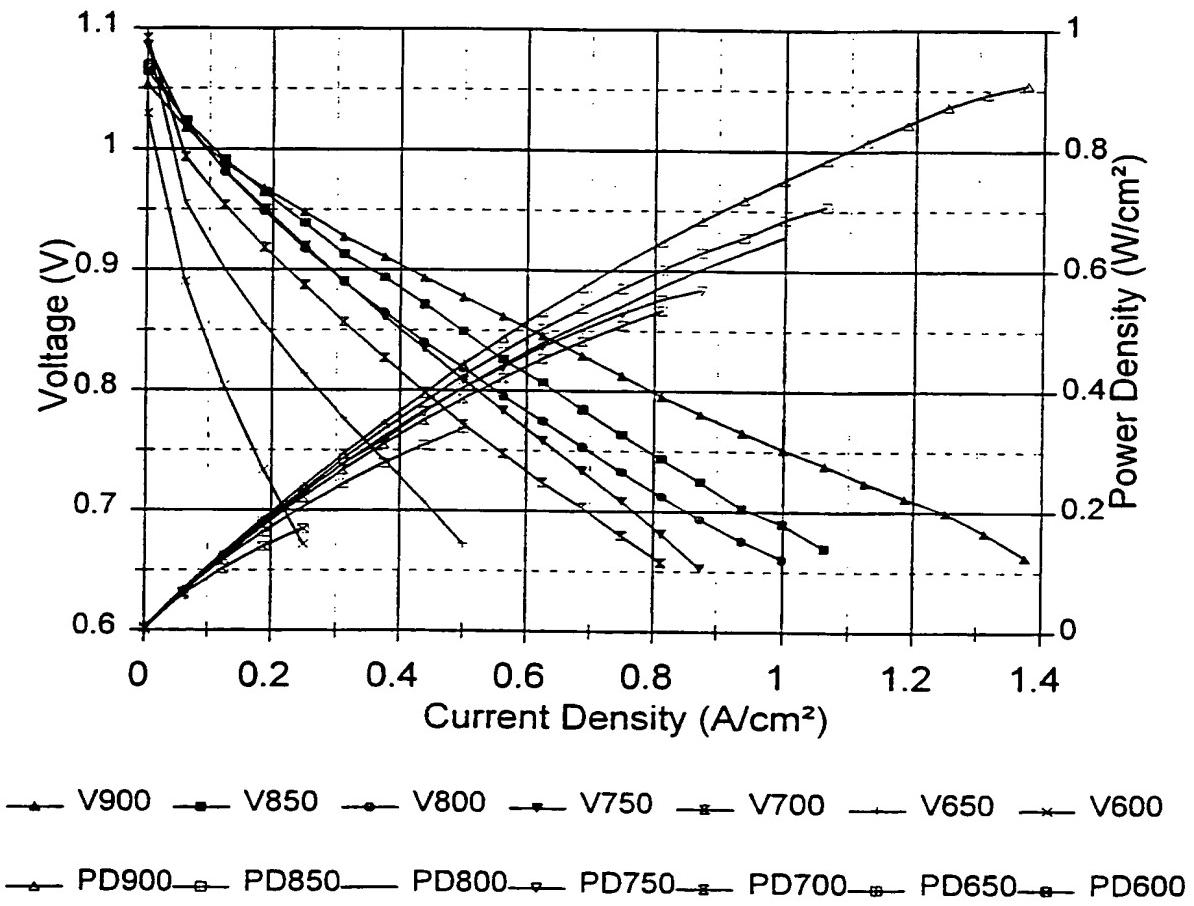




FIGURE 4

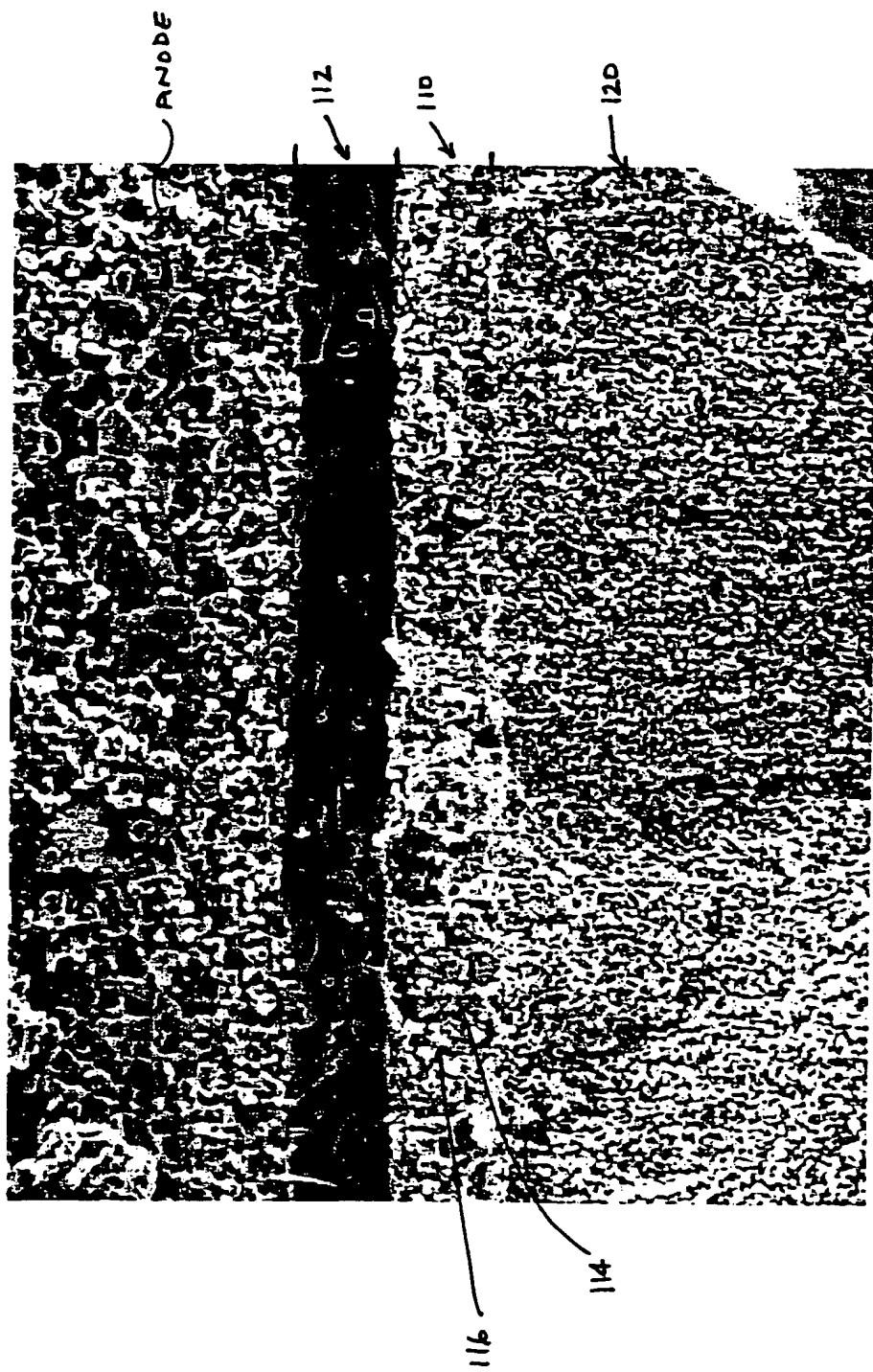


FIGURE 5

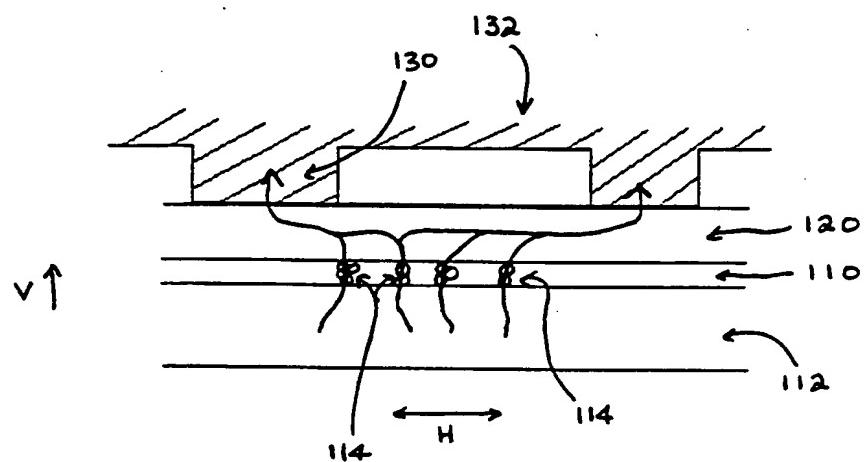


FIG. 6

FIGURE 7

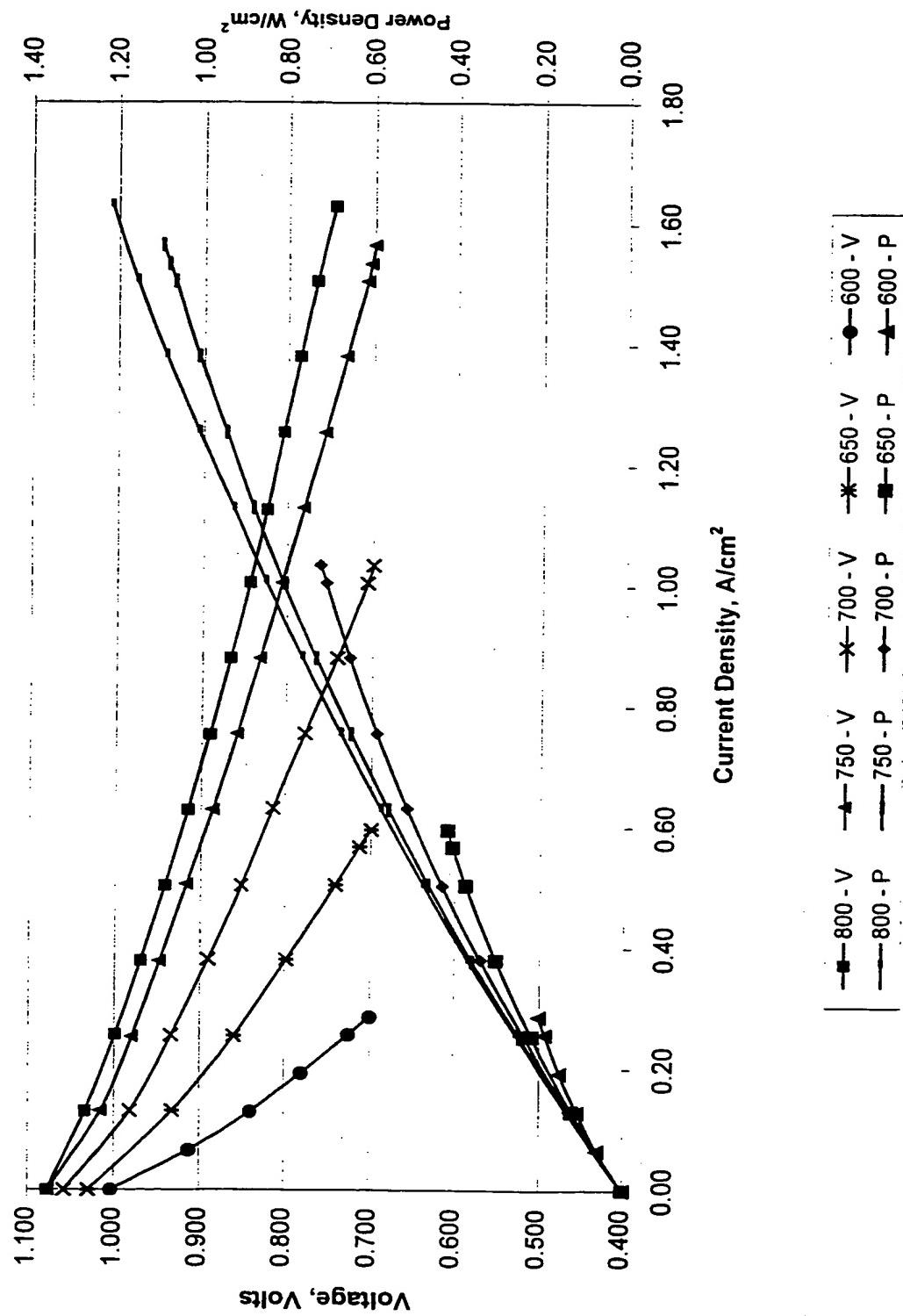
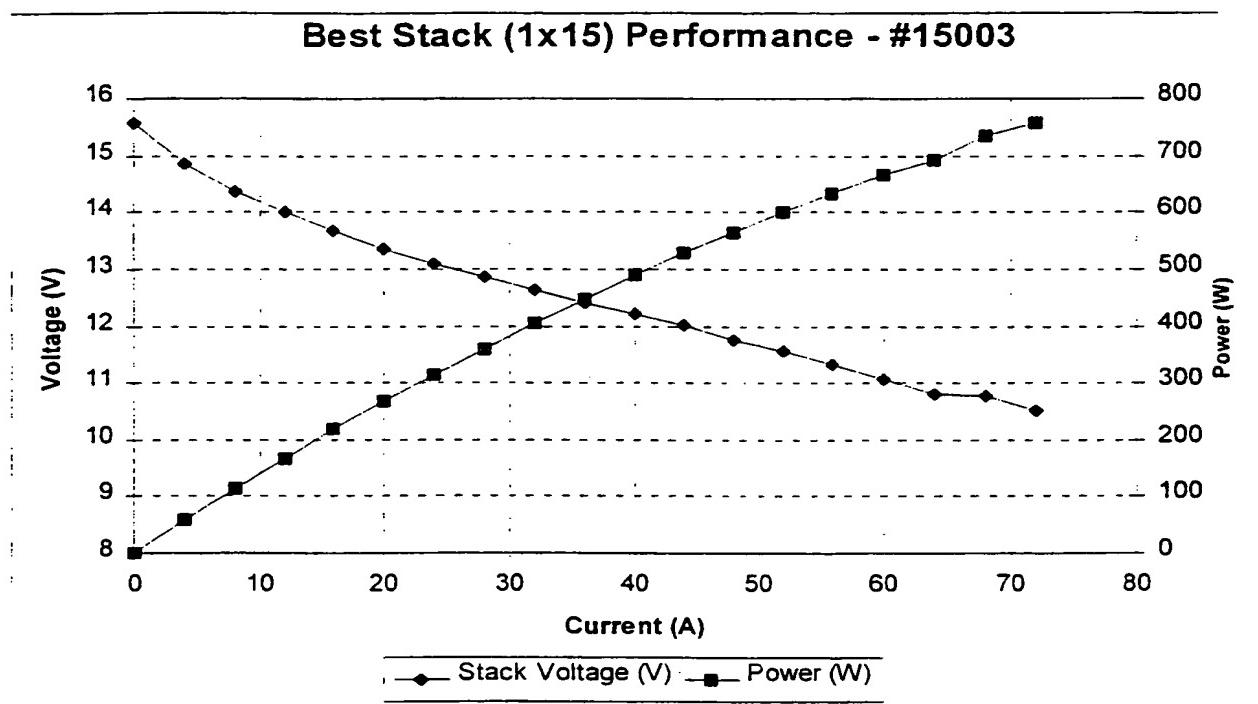


FIGURE 8



Alternative Cathode Loadings on 5 X 5 Cells  
1 lpm Hydrogen & 2 lpm Air

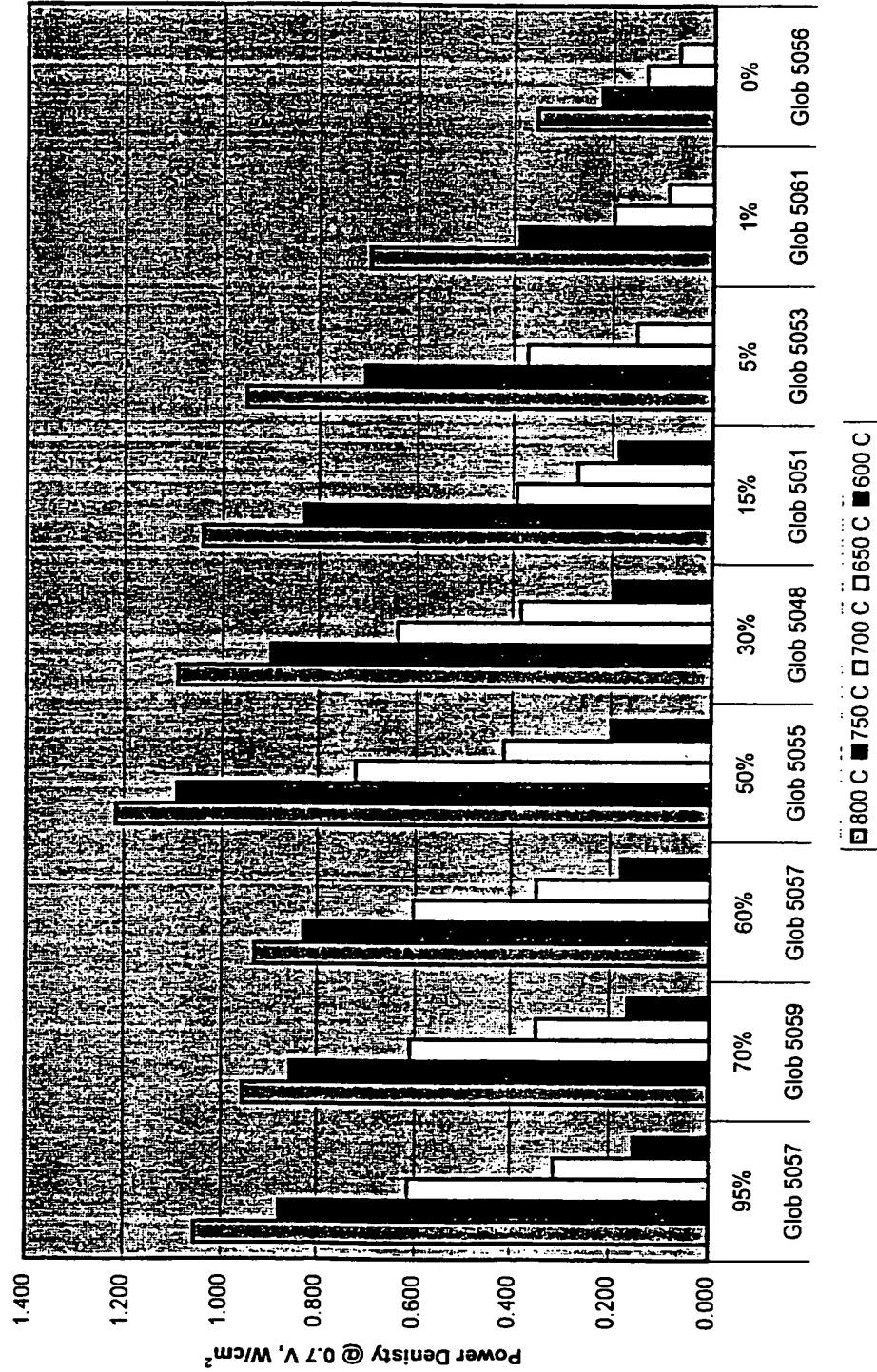


FIGURE 9

## INTERNATIONAL SEARCH REPORT

Inte	onal Application No
PCT/CA 00/01166	

**A. CLASSIFICATION OF SUBJECT MATTER**  
 IPC 7 H01M8/12 H01M4/86 G01N27/56

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
 IPC 7 H01M B01D G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, PAJ, WPI Data, CHEM ABS Data, INSPEC

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	PATENT ABSTRACTS OF JAPAN vol. 1998, no. 05, 30 April 1998 (1998-04-30) -& JP 10 021930 A (TOKYO GAS CO LTD), 23 January 1998 (1998-01-23) abstract ---	1,2,7,8, 18
X	PATENT ABSTRACTS OF JAPAN vol. 1998, no. 05, 30 April 1998 (1998-04-30) -& JP 10 021931 A (KYOCERA CORP;TOKYO GAS CO LTD), 23 January 1998 (1998-01-23) abstract --- -/-	1-3,6-8, 33,35,37

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

## ° Special categories of cited documents :

- \*A\* document defining the general state of the art which is not considered to be of particular relevance
- \*E\* earlier document but published on or after the international filing date
- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

- \*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- \*X\* document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- \*Y\* document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- \*&\* document member of the same patent family

Date of the actual completion of the international search

20 February 2001

Date of mailing of the international search report

06/03/2001

Name and mailing address of the ISA  
 European Patent Office, P.B. 5818 Patentlaan 2  
 NL - 2280 HV Rijswijk  
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
 Fax: (+31-70) 340-3016

Authorized officer

D'hondt, J

## INTERNATIONAL SEARCH REPORT

Inte  
Final Application No  
PCT/CA 00/01166

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	PATENT ABSTRACTS OF JAPAN vol. 016, no. 513 (E-1283), 22 October 1992 (1992-10-22) -& JP 04 192261 A (NGK INSULATORS LTD), 10 July 1992 (1992-07-10) abstract ---	1,2,6,18
X	DATABASE WPI Section Ch, Week 199107 Derwent Publications Ltd., London, GB; Class E36, AN 1991-046784 XP002160569 -& JP 02 310459 A (NGK SPARK PLUG CO LTD), 26 December 1990 (1990-12-26) abstract -& PATENT ABSTRACTS OF JAPAN vol. 015, no. 099, 11 March 1991 (1991-03-11) & JP 02 310459 A (NGK SPARK PLUG CO LTD), 26 December 1990 (1990-12-26) abstract ---	1-3
X	PATENT ABSTRACTS OF JAPAN vol. 016, no. 246 (E-1212), 5 June 1992 (1992-06-05) -& JP 04 051462 A (YUASA BATTERY CO LTD), 19 February 1992 (1992-02-19) abstract -& DATABASE CHEMABS 'Online! CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; OKUYAMA, RYOICHI: "Manufacture of electrodes and electrochemical devices" Database accession no. 117-30547 XP002160568 abstract ---	1
X	US 4 940 528 A (OKI SHUICHIRO ET AL) 10 July 1990 (1990-07-10) claims 1,6; table 1 ---	1-4,18, 19
P,X	WO 99 57779 A (BERKEL FRANCISCUS P F ;JONG JAN PETER DE (NL); SCHIPPER GERARDUS S) 11 November 1999 (1999-11-11) claims 1-3,5-10; figure 2 page 4, last paragraph -page 5, paragraph 1 page 2, line 28 page 2, line 31 - line 33 page 4, line 25 -page 31 ---	1-3,6-8, 33-35, 37,40
	-/-	

## INTERNATIONAL SEARCH REPORT

International Application No
PCT/CA 00/01166

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	DATABASE WPI Section Ch, Week 199312 Derwent Publications Ltd., London, GB; Class L03, AN 1993-097396 XP002160570 -& JP 05 041218 A (OSAKA GAS CO LTD), 19 February 1993 (1993-02-19) abstract ----	1
A	US 4 283 441 A (HAECKER WOLF-DIETER ET AL) 11 August 1981 (1981-08-11) example 1 ----	1
A	EP 0 439 938 A (WESTINGHOUSE ELECTRIC CORP) 7 August 1991 (1991-08-07) claims 1,2 ----	17
A	US 5 527 635 A (OKUO TAKAYASU ET AL) 18 June 1996 (1996-06-18) claims 1,2 ----	15
A	US 4 280 890 A (FRIESE KARL-HERMANN ET AL) 28 July 1981 (1981-07-28) column 4, line 35 - line 37; claim 1 ----	15
A	GB 2 298 955 A (BRITISH NUCLEAR FUELS PLC) 18 September 1996 (1996-09-18) claims 1,7,8 -----	15

**INTERNATIONAL SEARCH REPORT**

Information on patent family members

Inte...nal Application No

PCT/CA 00/01166

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
JP 10021930 A	23-01-1998	NONE		
JP 10021931 A	23-01-1998	NONE		
JP 04192261 A	10-07-1992	NONE		
JP 2310459 A	26-12-1990	JP 2790852 B		27-08-1998
JP 04051462 A	19-02-1992	NONE		
US 4940528 A	10-07-1990	JP 1184457 A		24-07-1989
WO 9957779 A	11-11-1999	NL 1009060 C AU 3852699 A		05-11-1999 23-11-1999
JP 5041218 A	19-02-1993	JP 3062632 B		12-07-2000
US 4283441 A	11-08-1981	DE 2852638 A CA 1123969 A JP 55078246 A SE 443876 B SE 7910030 A		19-06-1980 18-05-1982 12-06-1980 10-03-1986 07-06-1980
EP 0439938 A	07-08-1991	US 4971830 A CA 2034707 A DE 69011839 D DE 69011839 T ES 2058824 T JP 3058702 B JP 4215254 A KR 218599 B NO 910312 A		20-11-1990 02-08-1991 29-09-1994 27-04-1995 01-11-1994 04-07-2000 06-08-1992 01-09-1999 02-08-1991
US 5527635 A	18-06-1996	JP 2636157 B JP 7161361 A GB 2284599 A, B		30-07-1997 23-06-1995 14-06-1995
US 4280890 A	28-07-1981	DE 2911042 A JP 1440485 C JP 55128149 A JP 62049582 B		02-10-1980 30-05-1988 03-10-1980 20-10-1987
GB 2298955 A	18-09-1996	AU 705607 B AU 5114496 A CA 2215165 A CN 1183855 A EP 0815608 A WO 9628856 A JP 11502052 T NO 974249 A NZ 304057 A US 6013386 A ZA 9602172 A		27-05-1999 02-10-1996 19-09-1996 03-06-1998 07-01-1998 19-09-1996 16-02-1999 10-10-1997 29-04-1999 11-01-2000 31-07-1996

## REVISED VERSION

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
19 April 2001 (19.04.2001)

PCT

(10) International Publication Number  
WO 01/28024 A1

(51) International Patent Classification<sup>7</sup>: H01M 8/12. 4/86, G01N 27/407

(72) Inventors: GHOSH, Debabrata; 4852 - 52 Street S.E., Calgary, Alberta T2B 3R2 (CA). MARTELL, Frank; 4852 - 52 Street S.E., Calgary, Alberta T2B 3R2 (CA). TANG, Zheng; 4852 - 52 Street S.E., Calgary, Alberta T2B 3R2 (CA).

(74) Agent: BENNETT JONES LLP; 1000 Atco Centre, 10035 - 105 Street, Edmonton, Alberta TSJ 3T2 (CA).



patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

**Published:**

— with international search report

**(88) Date of publication of the revised international search report:**

11 October 2001

**(15) Information about Correction:**

see PCT Gazette No. 41/2001 of 11 October 2001, Section II

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

A. CLASSIFICATION OF SUBJECT MATTER  
 IPC 7 H01M8/12 H01M4/86 G01N27/407

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
 IPC 7 H01M B01D G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, PAJ, WPI Data, CHEM ABS Data, INSPEC

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	PATENT ABSTRACTS OF JAPAN vol. 1998, no. 05, 30 April 1998 (1998-04-30) -& JP 10 021930 A (TOKYO GAS CO LTD), 23 January 1998 (1998-01-23) abstract --- X	1,2,7,8, 18
X	PATENT ABSTRACTS OF JAPAN vol. 1998, no. 05, 30 April 1998 (1998-04-30) -& JP 10 021931 A (KYOCERA CORP; TOKYO GAS CO LTD), 23 January 1998 (1998-01-23) abstract --- -/-	1-3,6-8, 33,35,37

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

## \* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

20.02.01

Date of mailing of the international search report

23.07.01

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl.  
Fax: (+31-70) 340-3016

Authorized officer

D'hondt, J

## INTERNATIONAL SEARCH REPORT

International Application No

PCT/CA 00/01166

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	PATENT ABSTRACTS OF JAPAN vol. 016, no. 513 (E-1283), 22 October 1992 (1992-10-22) -& JP 04 192261 A (NGK INSULATORS LTD), 10 July 1992 (1992-07-10) abstract ---	1,2,6,18
X	DATABASE WPI Section Ch, Week 199107 Derwent Publications Ltd., London, GB; Class E36, AN 1991-046784 XP002160569 -& JP 02 310459 A (NGK SPARK PLUG CO LTD), 26 December 1990 (1990-12-26) abstract -& PATENT ABSTRACTS OF JAPAN vol. 015, no. 099, 11 March 1991 (1991-03-11) & JP 02 310459 A (NGK SPARK PLUG CO LTD), 26 December 1990 (1990-12-26) abstract ---	1-3
X	PATENT ABSTRACTS OF JAPAN vol. 016, no. 246 (E-1212), 5 June 1992 (1992-06-05) -& JP 04 051462 A (YUASA BATTERY CO LTD), 19 February 1992 (1992-02-19) abstract -& DATABASE CHEMABS [Online] CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; OKUYAMA, RYOICHI: "Manufacture of electrodes and electrochemical devices" Database accession no. 117-30547 XP002160568 abstract ---	1
X	US 4 940 528 A (OKI SHUICHIRO ET AL) 10 July 1990 (1990-07-10) claims 1,6; table 1 ---	1-4,18, 19
P,X	WO 99 57779 A (BERKEL FRANCISCUS P F ;JONG JAN PETER DE (NL); SCHIPPER GERARDUS S) 11 November 1999 (1999-11-11) claims 1-3,5-10; figure 2 page 4, last paragraph -page 5, paragraph 1 page 2, line 28 page 2, line 31 - line 33 page 4, line 25 -page 31 ---	1-3,6-8, 33-35, 37,40

## INTERNATIONAL SEARCH REPORT

International Application No

PCT/CA 00/01166

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	DATABASE WPI Section Ch, Week 199312 Derwent Publications Ltd., London, GB; Class L03, AN 1993-097396 XP002160570 -& JP 05 041218 A (OSAKA GAS CO LTD), 19 February 1993 (1993-02-19) abstract ---	1
A	US 4 283 441 A (HAECKER WOLF-DIETER ET AL) 11 August 1981 (1981-08-11) example 1 ---	1
A	EP 0 439 938 A (WESTINGHOUSE ELECTRIC CORP) 7 August 1991 (1991-08-07) claims 1,2 ---	17
A	US 5 527 635 A (OKUO TAKAYASU ET AL) 18 June 1996 (1996-06-18) claims 1,2 ---	15
A	US 4 280 890 A (FRIESE KARL-HERMANN ET AL) 28 July 1981 (1981-07-28) column 4, line 35 - line 37; claim 1 ---	15
A	GB 2 298 955 A (BRITISH NUCLEAR FUELS PLC) 18 September 1996 (1996-09-18) claims 1,7,8 -----	15

## INTERNATIONAL SEARCH REPORT

Information on patent family members

Inte-  
nal Application No

PCT/CA 00/01166

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
JP 10021930	A	23-01-1998	NONE		
JP 10021931	A	23-01-1998	NONE		
JP 04192261	A	10-07-1992	NONE		
JP 2310459	A	26-12-1990	JP	2790852 B	27-08-1998
JP 04051462	A	19-02-1992	NONE		
US 4940528	A	10-07-1990	JP	1184457 A	24-07-1989
WO 9957779	A	11-11-1999	NL	1009060 C	05-11-1999
			AU	3852699 A	23-11-1999
JP 5041218	A	19-02-1993	JP	3062632 B	12-07-2000
US 4283441	A	11-08-1981	DE	2852638 A	19-06-1980
			CA	1123969 A	18-05-1982
			JP	55078246 A	12-06-1980
			SE	443876 B	10-03-1986
			SE	7910030 A	07-06-1980
EP 0439938	A	07-08-1991	US	4971830 A	20-11-1990
			CA	2034707 A	02-08-1991
			DE	69011839 D	29-09-1994
			DE	69011839 T	27-04-1995
			ES	2058824 T	01-11-1994
			JP	3058702 B	04-07-2000
			JP	4215254 A	06-08-1992
			KR	218599 B	01-09-1999
			NO	910312 A	02-08-1991
US 5527635	A	18-06-1996	JP	2636157 B	30-07-1997
			JP	7161361 A	23-06-1995
			GB	2284599 A, B	14-06-1995
US 4280890	A	28-07-1981	DE	2911042 A	02-10-1980
			JP	1440485 C	30-05-1988
			JP	55128149 A	03-10-1980
			JP	62049582 B	20-10-1987
GB 2298955	A	18-09-1996	AU	705607 B	27-05-1999
			AU	5114496 A	02-10-1996
			CA	2215165 A	19-09-1996
			CN	1183855 A	03-06-1998
			EP	0815608 A	07-01-1998
			WO	9628856 A	19-09-1996
			JP	11502052 T	16-02-1999
			NO	974249 A	10-10-1997
			NZ	304057 A	29-04-1999
			US	6013386 A	11-01-2000
			ZA	9602172 A	31-07-1996